DRUG: Enoxaparin (Lovenox) or Low Molecular Weight Heparin

THERAPY: Home (Outpatient) Treatement of Deep Vein Thrombosis
Prophylaxis of Venous Thromboembolism
Perioperative Bridging for Chronic Oral Anticoagulation

REQUIRED BASELINE INFORMATION:

CBC with platelets
serum creatinine
patient weight (kg)
blood pressure

QUANTITY/DURATION OF THERAPY:

No prior authorization required for initial 5 days (10 dosages) of therapy. Prior authorization required for refill per established criteria.

EXCLUSION:

- Hypersensitivity to heparin or low molecular weight heparin products
- Hypersensitivity to pork products
- (+) in vitro test for anti-platelet antibody associated with thrombocytopenia
- history of heparin-induced thrombocytopenia
- platelet count less than 150,000
- active major bleeding
- surgery or acute bleeding event within 30 days
- history of intracranial hemorrhage or neoplasm
- end-stage renal disease requiring dialysis
- calculated creatinine clearance (CrCL) less than 30 mL/min
- patient weight less than 45kg (female) or 57kg (male)
- pregnant patient with mechanical prosthetic heart valve replacement
- increased hemorrhagic risk including (patient-specific)
 - congenital or acquired bleeding disorder
 - active ulceration and angiodysplastic gastrointestinal disease
 - □ hemorrhagic cerebral vascular accident
 - □ recent brain, spinal or ophthalmological surgery
 - □ bacterial endocarditis
 - □ severe uncontrolled hypertension
- Suspected or proven pulmonary embolism (PE)
- History of failed heparin therapy
- History of non-compliance with medical care and/or medications
- Communication barriers (no telephone, rural location, transient)
- Greater than 15 minutes from emergency services
- Current history of alcohol abuse
- Known risk of falls

INCLUSION:

	ute Deep Vein Thrombosis documented DVT by objective measures enrollment in established home health care program or resident at care facility established medical care for follow-up or resident at care facility concurrent oral anticoagulation therapy (warfarin) initiated Day #1
Pr	ophylaxis of VTE
	status post qualifying procedure total knee arthroplasty (TKA) total hip arthroplasty (THA) lower extremity trauma (pelvic and/or acetabular fracture) + non-weight bearing spinal cord injury abdominal surgery + non-ambulatory
	enrollment in established home health care program or residency at care facility concurrent oral anticoagulation therapy (warfarin) initiated Day #1 (if prophylaxis indicated beyond 96 hours)
Pe	rioperative Bridging for Chronic Oral Anticoagulation
	qualifying high-risk indication necessitating perioperative bridging o mechanical mitral heart valve (high-risk) o mechanical aortic heart valve (intermediate-risk) o mechanical mitral or aortic cage-ball valve (high-risk) o antiphospholipid syndrome (high-risk) o malignancy with associated thromboembolic risk (high-risk) o acute venous thromboembolic event within 90 days (high-risk) o acute arterial thromboembolic event within 30 days (high-risk) o acute pulmonary embolism within 90 days (high-risk) o atrial fibrillation with history of associated cerebral vascular accident (high-risk) qualifying invasive procedure with high risk of bleeding complications
	qualifying invasive procedure with high-risk of bleeding complications o colonoscopy with anticipated polypectomy (excluding screening colonoscopy) o central nervous system (CNS) surgery o vascular surgery o major thoracic, abdominal or pelvic surgery o high bleeding risk patient-specific surgical procedure (documentation required) (excluding dental procedures, cutaneous surgery and cataract surgery)
	warfarin stopped 5 days before scheduled procedure date low molecular weight heparin (1mg/kg Q12hr) initiated 36 hours after last warfarin dose low molecular weight heparin stopped 12 – 24 hours prior to procedure re-initiate low molecular weight heparin + warfarin 24 hours after procedure
Mo Da	edical Directorte: